

## **ENGROSSED** SENATE BILL No. 153

DIGEST OF SB 153 (Updated March 30, 1999 8:42 pm - DI 77)

Citations Affected: IC 25-26; IC 27-8.

**Synopsis:** Insurance coverage of mail order and Internet pharmacies. Amends definition of "practice of pharmacy" to include certain acts, including counseling. Allows insurers to designate a mail order or Internet pharmacy to provide prescription drugs to an insured. Prohibits an insurer from requiring as a condition of coverage that an insured purchase prescription drugs from a designated mail order or Internet pharmacy. Prohibits an insurer from applying a financial penalty if an insured does not purchase prescription drugs from an in-network designated mail order or Internet pharmacy. Requires a mail order or Internet pharmacy to comply with the laws of the state of domicile and with the Indiana generic drug law.

Effective: July 1, 1999.

# Gard, Sipes, Lubbers

(HOUSE SPONSORS — HASLER, WHETSTONE)

January 6, 1999, read first time and referred to Committee on Health and Provider

Vices. February 25, 1999, amended, reported favorably — Do Pass. March 1, 1999, read second time, ordered engrossed. March 2, 1999, engrossed. March 3, 1999, read third time, passed. Yeas 48, nays 0.

HOUSE ACTION

March 8, 1999, read first time and referred to Committee on Public Health.

April 5, 1999, amended, reported — Do Pass.



First Regular Session 111th General Assembly (1999)

PRINTING CODE. Amendments: Whenever an existing statute (or a section of the Indiana Constitution) is being amended, the text of the existing provision will appear in this style type, additions will appear in this style type, and deletions will appear in this style type.

Additions: Whenever a new statutory provision is being enacted (or a new constitutional provision adopted), the text of the new provision will appear in **this style type**. Also, the word **NEW** will appear in that style type in the introductory clause of each SECTION that adds a new provision to the Indiana Code or the Indiana Constitution.

Conflict reconciliation: Text in a statute in *this style type* or *this style type* reconciles conflicts between statutes enacted by the 1998 General Assembly.

## ENGROSSED SENATE BILL No. 153

A BILL FOR AN ACT to amend the Indiana Code concerning pharmacies.

Be it enacted by the General Assembly of the State of Indiana:

1	SECTION 1. IC 25-26-13-2 IS AMENDED TO READ AS
2	FOLLOWS [EFFECTIVE JULY 1, 1999]: Sec. 2. As used in this
3	chapter:

"Board" means the Indiana board of pharmacy.

"Controlled drugs" are those drugs on schedules I through V of the Federal Controlled Substances Act or on schedules I through V of IC 35-48-2.

"Counseling" means effective communication between a pharmacist and a patient concerning the contents, drug to drug interactions, route, dosage, form, directions for use, precautions, and effective use of a drug or device to improve the therapeutic outcome of the patient through the effective use of the drug or device.

"Dispensing" means issuing one (1) or more doses of a drug in a suitable container with appropriate labeling for subsequent administration to or use by a patient.

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1	"Drug" means:
2	(1) articles or substances recognized in the official United States
3	Pharmacopoeia, official National Formulary, official
4	Homeopathic Pharmacopoeia of the United States, or any
5	supplement to any of them;
6	(2) articles or substances intended for use in the diagnosis, cure,
7	mitigation, treatment, or prevention of disease in man or animals;
8	(3) articles other than food intended to affect the structure or any
9	function of the body of man or animals; or
10	(4) articles intended for use as a component of any article
11	specified in subdivisions (1) through (3) and devices.
12	"Drug order" means a written order in a hospital or other health care
13	institution for an ultimate user for any drug or device, issued and
14	signed by a practitioner, or an order transmitted by other means of
15	communication from a practitioner, which is immediately reduced to
16	writing by the pharmacist, registered nurse, or other licensed health
17	care practitioner authorized by the hospital or institution. The order
18	shall contain the name and bed number of the patient; the name and
19	strength or size of the drug or device; unless specified by individual
20	institution policy or guideline, the amount to be dispensed either in
21	quantity or days; adequate directions for the proper use of the drug or
22	device when it is administered to the patient; and the name of the
23	prescriber.
24	"Drug regimen review" means the retrospective, concurrent,
25	and prospective review by a pharmacist of a patient's drug related
26	history that includes the following areas:
27	(1) Evaluation of prescriptions or drug orders and patient
28	records for drug allergies, rational therapy contradictions,
29	appropriate dose and route of administration, appropriate
30	directions for use, or duplicative therapies.
31	(2) Evaluation of prescriptions or drug orders and patient
32	records for drug-drug, drug-food, drug-disease, and
33	drug-clinical laboratory interactions.
34	(3) Evaluation of prescriptions or drug orders and patient
35	records for adverse drug reactions.
36	(4) Evaluation of prescriptions or drug orders and patient
37	records for proper utilization and optimal therapeutic
38	outcomes.
39	"Drug utilization review" means a program designed to
40	measure and assess on a retrospective and prospective basis the
41	proper use of drugs.

"Device" means an instrument, apparatus, implement, machine,



1	contrivance, implant, invitro reagent, or other similar or related article
2	including any component part or accessory, which is:
3	(1) recognized in the official United States Pharmacopoeia,
4	official National Formulary, or any supplement to them;
5	(2) intended for use in the diagnosis of disease or other conditions
6	or the cure, mitigation, treatment, or prevention of disease in man
7	or other animals; or
8	(3) intended to affect the structure or any function of the body of
9	man or other animals and which does not achieve any of its
10	principal intended purpose through chemical action within or on
11	the body of man or other animals and which is not dependent
12	upon being metabolized for the achievement of any of its
13	principal intended purposes.
14	"Investigational or new drug" means any drug which is limited by
15	state or federal law to use under professional supervision of a
16	practitioner authorized by law to prescribe or administer such drug.
17	"Legend drug" has the meaning set forth in IC 16-18-2-199.
18	"License" and "permit" are interchangeable and mean a written
19	certificate from the Indiana board of pharmacy for the practice of
20	pharmacy or the operation of a pharmacy.
21	"Person" means any individual, partnership, copartnership, firm,
22	company, corporation, association, joint stock company, trust, estate,
23	or municipality, or a legal representative or agent, unless this chapter
24	expressly provides otherwise.
25	"Practitioner" means a physician licensed under IC 25-22.5, a
26	veterinarian licensed under IC 15-5-1.1, a dentist licensed under
27	IC 25-14, a podiatrist licensed under IC 25-29, or any other person
28	licensed by law to prescribe and administer legend drugs in this state.
29	"Pharmacist" means a person licensed under this chapter.
30	"Pharmacist extern" means a pharmacy student enrolled full-time in
31	an approved school of pharmacy and who is working in a school
32	sponsored, board approved program related to the practice of
33	pharmacy.
34	"Pharmacist intern" means a person who is working to secure
35	additional hours of practice and experience prior to making application
36	for a license to practice as a pharmacist.
37	"Pharmacy" means any facility, department, or other place where
38	prescriptions are filled or compounded and are sold, dispensed, offered,
39	or displayed for sale and which has as its principal purpose the

dispensing of drug and health supplies intended for the general health,

welfare, and safety of the public, without placing any other activity on

a more important level than the practice of pharmacy.

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1	"The practice of pharmacy" or "the practice of the profession of
2	pharmacy" or the practice of the "profession of pharmacy" means a
3	patient oriented health care profession in which pharmacists
4	interact and consult with patients and with other health care
5	professionals concerning drugs and devices used to enhance
6	patients' wellness, prevent illness, and optimize outcomes, by
7	accepting responsibility for performing or supervising the
8	following acts, services, and operations:
9	(1) The offering of or performing of those acts, service operations,
0	or transactions incidental to the interpretation, evaluation, and
.1	implementation of a prescription prescriptions or drug orders.
2	(2) The compounding, labeling, administering, dispensing, or
.3	selling of drugs and devices, including radioactive substances,
4	whether dispensed on under a practitioner's prescription or
.5	<b>drug order</b> , or sold or given directly to the ultimate consumer. or
6	(3) The proper and safe storage and distribution of drugs and
7	devices.
8	(4) The maintenance of proper records of the receipt, storage,
9	sale, and dispensing of drugs and devices.
20	(5) and the responsibility for Counseling, advising, and
21	educating patients, patients' caregivers, and health care
22	providers and professionals, as necessary, as to the contents,
23	therapeutic values, hazards, uses, significant problems, risks,
24	and appropriate manner of use of drugs or and devices.
25	(6) Assessing, recording, and reporting events related to the
26	use of drugs or devices.
27	(7) Obtaining and maintaining patient profiles, patient drug
28	histories relating to therapy, other pharmacy records, and
29	other patient health records.
80	(8) Monitoring, recording, and reporting drug therapy and
31	use.
32	(9) Performing drug evaluation, drug utilization review, and
33	drug regimen review.
34	(10) Participation in the selection, storage, and distribution of
35	drugs, dietary supplements, and devices.
86	(11) Participation in drug or drug related research.
37	(12) Provision of the professional acts, professional decisions,
88	and professional services necessary to maintain all areas of a
89	patient's pharmacy related care as specifically authorized
Ю	under this article.

"Prescription" means a written order or an order transmitted by

other means of communication from a practitioner to or for an

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1	ultimate user for any drug or device containing the name and address
2	of the patient, the name and strength or size of the drug or device, the
3	amount to be dispensed, adequate directions for the proper use of the
4	drug or device by the patient, and the name of the practitioner issued
5	and signed by a practitioner. or an order transmitted by other means of
6	communication from a practitioner and which is immediately reduced
7	to writing by the pharmacist.
8	"Record" means all papers, letters, memoranda, notes, prescriptions,
9	drug orders, invoices, statements, patient medication charts or files,
10	computerized records, or other written indicia, documents or objects
11	which are used in any way in connection with the purchase, sale, or
12	handling of any drug or device.
13	"Sale" means every sale and includes:
14	(1) manufacturing, processing, transporting, handling, packaging,
15	or any other production, preparation, or repackaging;
16	(2) exposure, offer, or any other proffer;
17	(3) holding, storing, or any other possession;
18	(4) dispensing, giving, delivering, or any other supplying; and
19	(5) applying, administering, or any other using.
20	SECTION 2. IC 27-8-27.6 IS ADDED TO THE INDIANA CODE
21	AS A <b>NEW</b> CHAPTER TO READ AS FOLLOWS [EFFECTIVE
22	JULY 1, 1999]:
23	Chapter 27.6. Insurer Designation of Mail Order and Internet
24	Pharmacies
25	Sec. 1. (a) This chapter applies to every:
26	(1) policy of accident and sickness insurance (as defined in
27	IC 27-8-5-1), whether written on an individual basis, a group
28	basis, a franchise basis, or a blanket basis;
29	(2) group contract (as defined in IC 27-13-1-16) or individual
30	contract (as defined in IC 27-13-1-21) through which a health
31	maintenance organization furnishes health care services;
32	(3) health care plan of a state or local governmental entity
33	that provides coverage for health care services on a
34	self-insurance basis; and
35	(4) employee welfare benefit plan (as defined in 29 U.S.C.
36	1002) that is self-funded;
37	that is issued, delivered, executed, or renewed in Indiana on or
38	after July 1, 2000.

(b) This chapter does not apply to a policy or contract with a federal governmental entity to insure or administer the Medicare+Choice plan or a plan that covers military or civilian federal employees and their dependents.



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1	Sec. 2. As used in this chapter, "insurer" means a company, a	
2	firm, a partnership, an entity, an association, an order, a society,	
3	or a system:	
4	(1) making any of the kinds of insurance;	
5	(2) entering into any of the kinds of contracts; or	
6	(3) providing any of the coverage;	
7	described in section 1 of this chapter.	
8	Sec. 3. As used in this chapter, "insured" means an individual	
9	who is entitled to coverage under any policy, contract, or plan	
10	described in section 1 of this chapter.	
11	Sec. 4. As used in this chapter, "mail order or Internet based	
12	pharmacy" means a pharmacy that is located in Indiana or is a	
13	nonresident pharmacy (as defined in IC 25-26-17-2) that dispenses	
14	prescription drugs:	
15	(1) through the United States Postal Service or other delivery	
16	services; or	
17	(2) after receiving a request for prescription drugs through	
18	the Internet;	
19	to patients in Indiana.	
20	Sec. 5. (a) An insurer that provides coverage for prescription	
21	drugs may designate a mail order or Internet based pharmacy to	
22	provide prescription drugs to an insured.	
23	(b) An insurer may not:	
24	(1) require an insured to obtain prescription drugs from a	
25	pharmacy designated under subsection (a) as a condition of	
26	coverage; or	
27	(2) impose an additional fee or other financial penalty upon an	
28	insured if the insured obtains prescription drugs from an	V
29	in-network pharmacy not designated in subsection (a) that	
30	agrees to provide all pharmaceutical services under the terms	
31	and conditions that apply to a pharmacy designated in	
32	subsection (a).	
33	Sec. 6. A mail order or Internet based pharmacy shall comply	
34	with the following:	
35	(1) The licensure laws of the state in which the mail order or	
36	Internet based pharmacy is domiciled.	
37	(2) The generic drug laws of Indiana under IC 16-42-22.	



### COMMITTEE REPORT

Mr. President: The Senate Committee on Health and Provider Services, to which was referred Senate Bill No. 153, has had the same under consideration and begs leave to report the same back to the Senate with the recommendation that said bill be AMENDED as follows:

Delete the title and insert the following:

A BILL FOR AN ACT to amend the Indiana Code concerning pharmacies.

Page 2, between lines 11 and 12, begin a new paragraph and insert:

- "Sec. 4. As used in this chapter, "mail order or Internet based pharmacy" means a pharmacy that is located in Indiana or is a nonresident pharmacy (as defined in IC 25-26-17-2) that dispenses prescription drugs:
  - (1) through the United States Postal Service or other delivery services; or
  - (2) after receiving a request for prescription drugs through the Internet;

to patients in Indiana.".

Page 2, line 12, delete "4" and insert "5".

Page 2, line 13, delete ":" and insert "a mail order or Internet based pharmacy".

Page 2, delete lines 14 through 19.

Page 2, run in lines 13 through 20.

Page 2, after line 30, begin a new paragraph and insert:

- "Sec. 6. A mail order or Internet based pharmacy shall comply with the following:
  - (1) The licensure laws of the state in which the mail order or Internet based pharmacy is domiciled.
  - (2) The generic drug laws of Indiana under IC 16-42-22.".

and when so amended that said bill do pass.

(Reference is to SB 153 as introduced.)

MILLER, Chairperson

Committee Vote: Yeas 8, Nays 2.

### COMMITTEE REPORT

Mr. Speaker: Your Committee on Public Health, to which was referred Senate Bill 153, has had the same under consideration and begs leave to report the same back to the House with the recommendation that said bill be amended as follows:

Page 1, between the enacting clause and line 1, begin a new paragraph and insert:

"SECTION 1. IC 25-26-13-2 IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 1999]: Sec. 2. As used in this chapter:

"Board" means the Indiana board of pharmacy.

"Controlled drugs" are those drugs on schedules I through V of the Federal Controlled Substances Act or on schedules I through V of IC 35-48-2.

"Counseling" means effective communication between a pharmacist and a patient concerning the contents, drug to drug interactions, route, dosage, form, directions for use, precautions, and effective use of a drug or device to improve the therapeutic outcome of the patient through the effective use of the drug or device.

"Dispensing" means issuing one (1) or more doses of a drug in a suitable container with appropriate labeling for subsequent administration to or use by a patient.

"Drug" means:

- (1) articles or substances recognized in the official United States Pharmacopoeia, official National Formulary, official Homeopathic Pharmacopoeia of the United States, or any supplement to any of them;
- (2) articles or substances intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in man or animals;
- (3) articles other than food intended to affect the structure or any function of the body of man or animals; or
- (4) articles intended for use as a component of any article specified in subdivisions (1) through (3) and devices.

"Drug order" means a written order in a hospital or other health care institution for an ultimate user for any drug or device, issued and signed by a practitioner, or an order transmitted by other means of communication from a practitioner, which is immediately reduced to writing by the pharmacist, registered nurse, or other licensed health care practitioner authorized by the hospital or institution. The order shall contain the name and bed number of the patient; the name and strength or size of the drug or device; unless specified by individual

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С р institution policy or guideline, the amount to be dispensed either in quantity or days; adequate directions for the proper use of the drug or device when it is administered to the patient; and the name of the prescriber.

"Drug regimen review" means the retrospective, concurrent, and prospective review by a pharmacist of a patient's drug related history that includes the following areas:

- (1) Evaluation of prescriptions or drug orders and patient records for drug allergies, rational therapy contradictions, appropriate dose and route of administration, appropriate directions for use, or duplicative therapies.
- (2) Evaluation of prescriptions or drug orders and patient records for drug-drug, drug-food, drug-disease, and drug-clinical laboratory interactions.
- (3) Evaluation of prescriptions or drug orders and patient records for adverse drug reactions.
- (4) Evaluation of prescriptions or drug orders and patient records for proper utilization and optimal therapeutic outcomes.

"Drug utilization review" means a program designed to measure and assess on a retrospective and prospective basis the proper use of drugs.

"Device" means an instrument, apparatus, implement, machine, contrivance, implant, invitro reagent, or other similar or related article including any component part or accessory, which is:

- (1) recognized in the official United States Pharmacopoeia, official National Formulary, or any supplement to them;
- (2) intended for use in the diagnosis of disease or other conditions or the cure, mitigation, treatment, or prevention of disease in man or other animals; or
- (3) intended to affect the structure or any function of the body of man or other animals and which does not achieve any of its principal intended purpose through chemical action within or on the body of man or other animals and which is not dependent upon being metabolized for the achievement of any of its principal intended purposes.

"Investigational or new drug" means any drug which is limited by state or federal law to use under professional supervision of a practitioner authorized by law to prescribe or administer such drug.

"Legend drug" has the meaning set forth in IC 16-18-2-199.

"License" and "permit" are interchangeable and mean a written certificate from the Indiana board of pharmacy for the practice of

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pharmacy or the operation of a pharmacy.

"Person" means any individual, partnership, copartnership, firm, company, corporation, association, joint stock company, trust, estate, or municipality, or a legal representative or agent, unless this chapter expressly provides otherwise.

"Practitioner" means a physician licensed under IC 25-22.5, a veterinarian licensed under IC 15-5-1.1, a dentist licensed under IC 25-14, a podiatrist licensed under IC 25-29, or any other person licensed by law to prescribe and administer legend drugs in this state.

"Pharmacist" means a person licensed under this chapter.

"Pharmacist extern" means a pharmacy student enrolled full-time in an approved school of pharmacy and who is working in a school sponsored, board approved program related to the practice of pharmacy.

"Pharmacist intern" means a person who is working to secure additional hours of practice and experience prior to making application for a license to practice as a pharmacist.

"Pharmacy" means any facility, department, or other place where prescriptions are filled or compounded and are sold, dispensed, offered, or displayed for sale and which has as its principal purpose the dispensing of drug and health supplies intended for the general health, welfare, and safety of the public, without placing any other activity on a more important level than the practice of pharmacy.

"The practice of pharmacy" or "the practice of the profession of pharmacy" or the practice of the "profession of pharmacy" means a patient oriented health care profession in which pharmacists interact and consult with patients and with other health care professionals concerning drugs and devices used to enhance patients' wellness, prevent illness, and optimize outcomes, by accepting responsibility for performing or supervising the following acts, services, and operations:

- (1) The offering **of** or performing of those acts, service operations, or transactions incidental to the interpretation, **evaluation**, **and implementation** of a prescription **prescriptions** or **drug** orders.
- (2) The compounding, **labeling**, administering, dispensing, or selling of drugs and devices, **including radioactive substances**, whether dispensed <del>on</del> **under a practitioner's** prescription **or drug order**, or sold or given directly to the ultimate consumer. <del>or</del>
- (3) The proper and safe storage and distribution of drugs and devices.
- (4) The maintenance of proper records of the receipt, storage, sale, and dispensing of drugs and devices.



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- (5) and the responsibility for Counseling, advising, and educating patients, patients' caregivers, and health care providers and professionals, as necessary, as to the contents, therapeutic values, hazards, uses, significant problems, risks, and appropriate manner of use of drugs or and devices.
- (6) Assessing, recording, and reporting events related to the use of drugs or devices.
- (7) Obtaining and maintaining patient profiles, patient drug histories relating to therapy, other pharmacy records, and other patient health records.
- (8) Monitoring, recording, and reporting drug therapy and use.
- (9) Performing drug evaluation, drug utilization review, and drug regimen review.
- (10) Participation in the selection, storage, and distribution of drugs, dietary supplements, and devices.
- (11) Participation in drug or drug related research.
- (12) Provision of the professional acts, professional decisions, and professional services necessary to maintain all areas of a patient's pharmacy related care as specifically authorized under this article.

"Prescription" means a written order **or an order transmitted by other means of communication from a practitioner** to or for an
ultimate user for any drug or device containing the name and address
of the patient, the name and strength or size of the drug or device, the
amount to be dispensed, adequate directions for the proper use of the
drug or device by the patient, and the name of the practitioner issued
and signed by a practitioner. <del>or an order transmitted by other means of
communication from a practitioner and which is immediately reduced
to writing by the pharmacist.</del>

"Record" means all papers, letters, memoranda, notes, prescriptions, drug orders, invoices, statements, patient medication charts or files, computerized records, or other written indicia, documents or objects which are used in any way in connection with the purchase, sale, or handling of any drug or device.

"Sale" means every sale and includes:

- (1) manufacturing, processing, transporting, handling, packaging, or any other production, preparation, or repackaging;
- (2) exposure, offer, or any other proffer;
- (3) holding, storing, or any other possession;
- (4) dispensing, giving, delivering, or any other supplying; and
- (5) applying, administering, or any other using.".



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Page 1, line 1, delete "27-8-27" and insert "27-8-27.6".

Page 1, line 3, begin a new paragraph beginning with "Chapter".

Page 1, line 3, delete "27." and insert "27.6.".

Page 1, line 5, after "1." insert "(a)".

Page 2, line 1, delete "1999" and insert "2000".

Page 2, between lines 1 and 2, begin a new paragraph and insert:

"(b) This chapter does not apply to a policy or contract with a federal governmental entity to insure or administer the Medicare+Choice plan or a plan that covers military or civilian federal employees and their dependents.".

Page 2, line 29 delete "a" and insert "an in-network".

Page 2, line 31, after "provide" insert "all".

Renumber all SECTIONS consecutively.

and when so amended that said bill do pass.

(Reference is to SB 153 as printed February 26, 1999.)

BROWN C, Chair

Committee Vote: yeas 13, nays 0.

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